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Characterization of PMDI/HOPG Interfaces by SERS
Using a Silver Overlayer Configuration

by

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Presented

at

13th Annual Meeting of The Adhesion Society
Savannah, GA
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19. ABSTRACT (Continue on reverse if necessary and identify by block number) Surface-enhanced Raman scattering (SERS) was used for characterization of model fiber/polyimide interfaces. Pyromellitic diimide (PMDI) was used as a model polyimide while highly oriented pyrolytic graphite (HOPG) was viewed as a model graphite fiber. Samples for SERS were prepared by depositing films of PMDI onto HOPG and then depositing silver island films on top of the PMDI. When the PMDI films were relatively thick (85 Å), the SERS spectra were similar to normal Raman spectra of bulk PMDI. These spectra were characterized by the imide bands near 1775, 1748, 1368, and 659 cm^{-1} and by the benzene ring bands near 1633, 1200, 762, 578, and 520 cm^{-1} , respectively. For thin films of PMDI (about 15 Å), the SERS spectra were considerably different from normal Raman spectra of PMDI and SERS spectra of thick PMDI films on HOPG. The 1775 cm^{-1} band due to a C=O stretching vibration and the 762 cm^{-1} band related to a ring breathing mode were weak while the 1368 cm^{-1}					
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band due to a CNC axial stretching vibration and the 1200 cm^{-1} band due to a CNC axial stretching vibration and the 1200 cm^{-1} band due to a CX in-plane stretching mode were strong. Differences in the relative intensities of bands in the SERS spectra were attributed to orientation effects. It was concluded that PMDI films deposited on HOPG were actually bilayers in which the molecules adjacent to the surface were adsorbed with a vertical conformation in which the planes of the molecules were perpendicular to the surface and one imide group was in contact with the surface. Molecules farther away from the surface had a random orientation.

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